

**Listing of Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application.

Claim 1 (Currently amended): A pharmaceutical composition comprising one compound, which wherein the compound is a serotonin reuptake inhibitor, and another a second compound, which wherein the second compound is a H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist having an affinity for the H<sub>3</sub> receptor below 0.5 μM.

Claim 2 (Currently amended): A pharmaceutical composition comprising a compound that is both a H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist having an affinity for the H<sub>3</sub> receptor below 0.5 μM, and a serotonin reuptake inhibitor.

Claim 3 (Currently amended): A method of augmenting and/or providing faster onset of the therapeutic effect of a serotonin reuptake inhibitor, comprising administering to a patient in need thereof a therapeutically effective amount of a H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist having an affinity for the H<sub>3</sub> receptor below 0.5 μM.

Claim 4 (Currently amended): A method of treating depression or an affective disorder, comprising administering a therapeutically effective amount of a H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist having an affinity for the H<sub>3</sub> receptor below 0.5 μM to a patient being treated with a serotonin reuptake inhibitor and in need thereof.

Claim 5 (Previously presented): A method of treating depression or an affective disorder, comprising administering to a patient in need thereof a therapeutically effective amount of a pharmaceutical composition according to claim 1.

Claim 6 (Currently amended): The method of claim 5, wherein the compound serotonin reuptake inhibitor is a selective serotonin reuptake inhibitor.

Claim 7 (Currently amended): The method of claim 5, wherein the compound H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist is selective for the H<sub>3</sub> receptor.

Claim 8 (Currently amended): The method of claim 5, wherein the second compound is an antagonist or an inverse agonist at the H<sub>3</sub> receptor.

Claim 9 (Currently amended): The method of claim 5 [[8]], wherein the second compound is a H<sub>3</sub> receptor antagonist.

Claim 10 (Currently amended): The method of claim [[6]] 5, wherein the serotonin reuptake inhibitor is selected from citalopram, escitalopram, fluoxetine, sertraline, paroxetine, fluvoxamine, venlafaxine, dapoxetine, duloxetine, vilazodone, nefazodone, imipramine, femoxetine, or [[and]] clomipramine.

Claim 11 (Currently amended): The method of claim 5, wherein the H<sub>3</sub> receptor ligand H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist is selected from Thioperamide, Ciproxifan, Iodophenpropit, GR 175737, Iodoproxyfan, Proxifan, Perceptin, JB 98064, VUF 9153, A 304121, ABT923, ABT 834, A 923, A 320436, A 331440, A 349413, A 349821, A 417022, A 423579, A 424835, A 431404, AQ 0145, FUB 181, FUB 360, FUB 407, FUB 637, FUB 836, GR 168320, GSK 189254A, GSK 207040A, GT 2016, GT 2104, GT 2209, GT 2212, GT 2227, GT 2232, GT 2390, GT 2349, GT 2355, GT 2394, Imoproxifan, Impentamine, JNJ 5207852, NNC 0038 0000 1049, NNC 0038 0000 1202, SCH 50971, SCH 79687, UCL 1199, UCL 1283, UCL 1390, UCL 1409, UCL 1860, UCL 1972, UCL 2065, UCL 2138, UCL 2173, UCL 2283, Verongamine, VUF 4163, VUF 5000, or [[and]] VUF 5182.

Claim 12 (Previously presented): The pharmaceutical composition according to claim 1, further comprising a pharmaceutically acceptable carrier or diluent.

Claim 13 (Currently amended): The pharmaceutical composition according to claim 1, wherein the serotonin reuptake inhibitor used is a selective serotonin reuptake inhibitor.

Claim 14 (Previously presented): The pharmaceutical composition according to claim 1, wherein the H<sub>3</sub> antagonist, inverse agonist or partial agonist is selective for the H<sub>3</sub> receptor.

Claim 15 (Currently amended): The pharmaceutical composition according to claim 1, wherein the second compound H<sub>3</sub> ligand is a H<sub>3</sub> receptor antagonist.

Claim 16 (Currently amended): The pharmaceutical composition according to claim 1, wherein the serotonin uptake inhibitor is selected from citalopram, escitalopram, fluoxetine, sertraline, paroxetine, fluvoxamine, venlafaxine, dapoxetine, duloxetine, vilazodone, nefazodone, imipramine, femoxetine, or [[and]] clomipramine.

Claim 17 (Currently amended): The pharmaceutical composition according to claim 1, wherein the H<sub>3</sub> ligand antagonist, inverse agonist or partial agonist is selected from from Thioperamide, Ciproxifan, Iodophenpropit, GR 175737, Iodoproxyfan, Proxifan, Perceptin, JB 98064, VUF 9153, A 304121, ABT923, ABT 834, A 923, A 320436, A 331440, A 349413, A 349821, A 417022, A 423579, A 424835, A 431404, AQ 0145, FUB 181, FUB 360, FUB 407, FUB 637, FUB 836, GR 168320, GSK 189254A, GSK 207040A, GT 2016, GT 2104, GT 2209, GT 2212, GT 2227, GT 2232, GT 2390, GT 2349, GT 2355, GT 2394, Imoproxifan, Impentamine, JNJ 5207852, NNC 0038 0000 1049, NNC 0038 0000 1202, SCH 50971, SCH 79687, UCL 1199, UCL 1283, UCL 1390, UCL 1409, UCL 1860, UCL 1972, UCL 2065, UCL 2138, UCL 2173, UCL 2283, Verongamine, VUF 4163, VUF 5000, or [[and]] VUF 5182.

Claim 18 (Currently amended): The method of claim 5, wherein the active ingredients serotonin uptake inhibitor and H<sub>3</sub> antagonist, inverse agonist or partial agonist are administered by simultaneous administration.

Claim 19 (Currently amended): The method of claim 5, wherein the active ingredients serotonin uptake inhibitor and H<sub>3</sub> antagonist, inverse agonist or partial agonist are administered in the same unit dosage form.

Claim 20 (Currently amended): The method of claim 5, wherein the ~~active ingredients serotonin uptake inhibitor and H<sub>3</sub> antagonist, inverse agonist or partial agonist~~ are administered by sequential administration.

Claim 21 (Currently amended): The method of claim 5, wherein the ~~active ingredients serotonin uptake inhibitor and H<sub>3</sub> antagonist, inverse agonist or partial agonist~~ are administered in discrete dosage forms.

Claim 22 (Currently amended): A method for identifying compounds useful for the treatment of depression or an affective disorder, comprising, in any order:

(a) measuring the ability of ~~test~~ compounds to inhibit serotonin reuptake and selecting the compounds that have an IC<sub>50</sub> value below 50 nM;

(b) measuring the affinity of ~~test selected~~ compounds to the H<sub>3</sub> receptor and further selecting the compounds that have an affinity for the H<sub>3</sub> receptor below 0.5  $\mu$ M,

and thereafter measuring the efficacy of the selected compounds at the H<sub>3</sub> receptor and further selecting the compounds which are antagonists, inverse agonists or partial agonists at the H<sub>3</sub> receptor.

Claim 23 (Previously presented): The method according to claim 22 wherein the compound has an affinity in step (b) of less than 50 nM.

Claim 24 (Previously presented): The method according to claim 23, wherein the compound has an affinity in step (b) of less than 10 nM.

Claim 25 (Currently amended): A compound that inhibits serotonin reuptake and has an IC<sub>50</sub> value below 50 nM; and has an affinity to the H<sub>3</sub> receptor below 0.5  $\mu$ M.

Claim 26 (Previously presented): A method of treating depression or an affective disorder, comprising administering to a patient in need thereof a therapeutically effective amount of a pharmaceutical composition according to claim 2.

Claim 27 (Currently amended): The method of claim 26, wherein the compound serotonin reuptake inhibitor is a selective serotonin reuptake inhibitor.

Claim 28 (Currently amended): The method of claim 26, wherein the compound H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist is selective for the H<sub>3</sub> receptor.

Claim 29 (Currently amended): The method of claim 26, wherein the compound H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist is an antagonist or an inverse agonist at the H<sub>3</sub> receptor.

Claim 30 (Currently amended): The method of claim 29, wherein the compound H<sub>3</sub> receptor antagonist, inverse agonist or partial agonist is a H<sub>3</sub> receptor antagonist.

Claim 31 (Previously presented): The pharmaceutical composition according to claim 2, further comprising a pharmaceutically acceptable carrier or diluent.

Claim 32 (Currently amended): The pharmaceutical composition according to claim 2, wherein the compound H<sub>3</sub> ligand is a H<sub>3</sub> receptor antagonist.

Claim 33 (Currently amended): The method of claim 4, wherein the affective disorder is selected from an anxiety disorder disorders, generalized anxiety disorder, panic anxiety, obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder, social anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual syndrome, a cognitive disorder disorders, an impulse control disorder disorders, attention deficit hyperactivity disorder, or drug abuse or any other disorder responsive to a serotonin reuptake inhibitor.

Claim 34 (Currently amended): The method of claim 33, wherein the eating disorder is ~~selected from~~, bulimia, anorexia, or [[and]] obesity.

Claim 35 (Currently amended): The method of claim 5, wherein the affective disorder is ~~selected from~~ an anxiety disorder disorders, generalized anxiety disorder, panic anxiety, obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder, social anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual syndrome, a cognitive disorder disorders, an impulse control disorder disorders, attention deficit hyperactivity disorder, or drug abuse ~~or any other disorder responsive to a serotonin reuptake inhibitor~~.

Claim 36 (Currently amended): The method of claim 35, wherein the eating disorder is ~~selected from~~, bulimia, anorexia, or [[and]] obesity.

Claim 37 (Currently amended): The method of claim 22, wherein the affective disorder is ~~selected from~~ an anxiety disorder disorders, generalized anxiety disorder, panic anxiety, obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder, social anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual syndrome, a cognitive disorder disorders, an impulse control disorder disorders, attention deficit hyperactivity disorder, or drug abuse ~~or any other disorder responsive to a serotonin reuptake inhibitor~~.

Claim 38 (Currently amended): The method of claim 37, wherein the eating disorder is ~~selected from~~, bulimia, anorexia, or [[and]] obesity.

Claim 39 (Currently amended): The method of claim 26, wherein the affective disorder is ~~selected from~~ an anxiety disorder disorders, generalized anxiety disorder, panic anxiety, obsessive compulsive disorder, acute stress disorder, post traumatic stress disorder, social anxiety disorder, an eating disorder disorders, a phobia phobias, dysthymia, premenstrual

syndrome, a cognitive disorder disorders, an impulse control disorder disorders, attention deficit hyperactivity disorder, or drug abuse or any other disorder responsive to a serotonin reuptake inhibitor.

Claim 40 (Currently amended): The method of claim 39, wherein the eating disorder is selected from, bulimia, anorexia, or [[and]] obesity.

Claim 41 (Previously presented): The compound of claim 25, wherein the compound has an affinity to the H<sub>3</sub> receptor of less than 50 nM.

Claim 42 (Previously presented): The compound of claim 41, wherein the compound has an affinity to the H<sub>3</sub> receptor of less than 10 nM.